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(54) A phenylalkylaminoethylsalicylamide, its preparation and pharmaceutical compositions containing it.

(57) The (R,R) optical isomer of labetalol, namely 5-[(R)-1-hydroxy-2-[(R)-1-methyl-3-phenylpropyl]amino]ethyl]salicylamide, and its pharmaceutically acceptable acid addition salts, in a state substantially free of the corresponding (R,S), (S,R) and (S,S) optical isomers, are novel compounds with favourable therapeutic properties in comparison with labetalol. They can be used in particular in the treatment of hypertension.

The novel (R,R) optical isomer and its pharmaceutically acceptable acid addition salts can be prepared by removal of the protecting groups from an N,O-protected (R,R) optical isomer or acid addition salt thereof, wherein the basic nitrogen atom and the phenolic hydroxy group are protected, preferably by hydrogenolytic removal of hydrogenolysable groups.

EP 0 009 702 A1

A PHENYLALKYLAMINOETHYLSALICYLAMIDE, ITS PREPARATION
AND PHARMACEUTICAL COMPOSITIONS CONTAINING IT.

This invention relates to a phenylalkylaminoethylsalicyl-
amide, its preparation and pharmaceutical compositions
5 containing it. More particularly it relates to one op-
tical isomer of labetalol, 5-[1-hydroxy-2-(1-methyl-3-
-phenylpropylamino)-ethyl]salicylamide.

The substance labetalol is known from British patent specifi-
cation 1,266,058 and U.S.P. 4,012,444. Its pharmacological
10 properties are discussed by Farmer et. al. in British Jour-
nal of Pharmacology, 45: 660-675 (1972), who designate it
AH5158; it is shown to block α - and β -adrenergic receptors,
suggesting that it would be useful in the treatment of
arrhythmia, hypertension and angina pectoris.

15 The unique pharmacological properties of labetalol and its
use as an antihypertensive agent are said to be largely a
function of the exquisite balance of its α - and β -blocking
activities. The file history of U.S.P. 4,012,444 indeed in-
dicates that slight changes in the chemical structure of
20 labetalol deleteriously affect this balance, and, even in
the few analogous compounds where the balance is retained,
the absolute potencies of these compounds are shown to be
too low for them to be useful antihypertensive agents. There-
fore, in the treatment of hypertension, labetalol is the

compound of choice among those disclosed in British patent specification 1,266,058 and U.S.P. 4,012,444.

Labetalol has two asymmetrically substituted carbon atoms and therefore can exist as two diastereoisomers and four optical isomers. Indeed, British patent specification 1,266,058 and U.S.P. 4,012,444 disclose that compounds such as labetalol have optically active forms, but give no example of an optically active form. These patent specifications teach that "the racemic mixtures may be resolved by conventional methods, for example by salt formation with an optically active acid, followed by fractional crystallization", but give no method of resolution. Example 14 of each specification does indeed describe the separation of labetalol into two diastereoisomers "1" and "2", using benzoic acid, but this is not an optical resolution. In British patent specifications 1,541,932 and 1,541,933, "isomer 1" is designated "diastereoisomer A" and is characterised as that diastereoisomer whose hydrochloride salt has the higher melting point. These two British patent specifications also disclose that diastereoisomer A is a valuable antiarrhythmic agent since it has strongly reduced β -adrenergic blocking activity and is therefore useful in the treatment of people who have suffered myocardial infarction.

We have now discovered that diastereoisomer A is composed of the (S,R) and (R,S) optical isomers of labetalol, whereas di-

astereoisomer B is composed of the (S,S) and (R,R) optical isomers. We have also surprisingly found that the novel (R,R) optical isomer of labetalol exhibits, in comparison with labetalol itself, both an unexpectedly high increase in β -adrenergic blocking potency and a decrease in α -adrenergic blocking potency. Thus, when the (R,R) optical isomer is compared with labetalol, the ratio of the β -adrenergic blocking potency to the α -adrenergic blocking potency is found to be greatly and unexpectedly increased. In particular, animal tests have indicated that the (R,R) optical isomer has about twelve times the β -blocking potency of labetalol, but only about one third of the α -blocking potency of labetalol. These properties could in no way have been predicted theoretically, especially as the β -blocking potency of diastereoisomer B is not significantly different from that of labetalol and the α -blocking potency of diastereoisomer B is half that of labetalol. Indeed, it is clear, when the activities of the four optical isomers of labetalol are compared, that the activities of the diastereoisomers A and B and indeed of labetalol itself cannot be calculated from the activities of their components. One can put this the other way around by saying that the α - and β -blocking activities of the four optical isomers of labetalol do not merely average to give the α - and β -blocking activities of labetalol and of its diastereoisomers A and B. Some of the activities are much greater than could ever have

been expected on a simple basis of mathematical proportions, in particular the high β -blocking activity of the (R,R) optical isomer: this activity is much higher than the β -blocking activity of diastereoisomer B so that antagonism evidently exists between the (S,S) and (R,R) optical isomers with respect to the β -blocking activity. This degree of antagonism could in no way have been foreseen. In the absence of this antagonism, the (R,R) optical isomer shows a balance of properties that make it the optical isomer of choice in the treatment of hypertension. In particular, the (R,R) optical isomer possesses potent antihypertensive activity and rapid onset of activity while substantially lacking the undesirable side-effects usually associated with α -blockade, e.g. postural hypotension.

The following Table shows the relationships between labetalol, its diastereoisomers A and B and the four pure optical isomers; below each compound are given its potencies as an α -blocking and then as a β -blocking agent, all relative to the values for labetalol (assigned values 1.0 for each blocking activity):

- 5 -

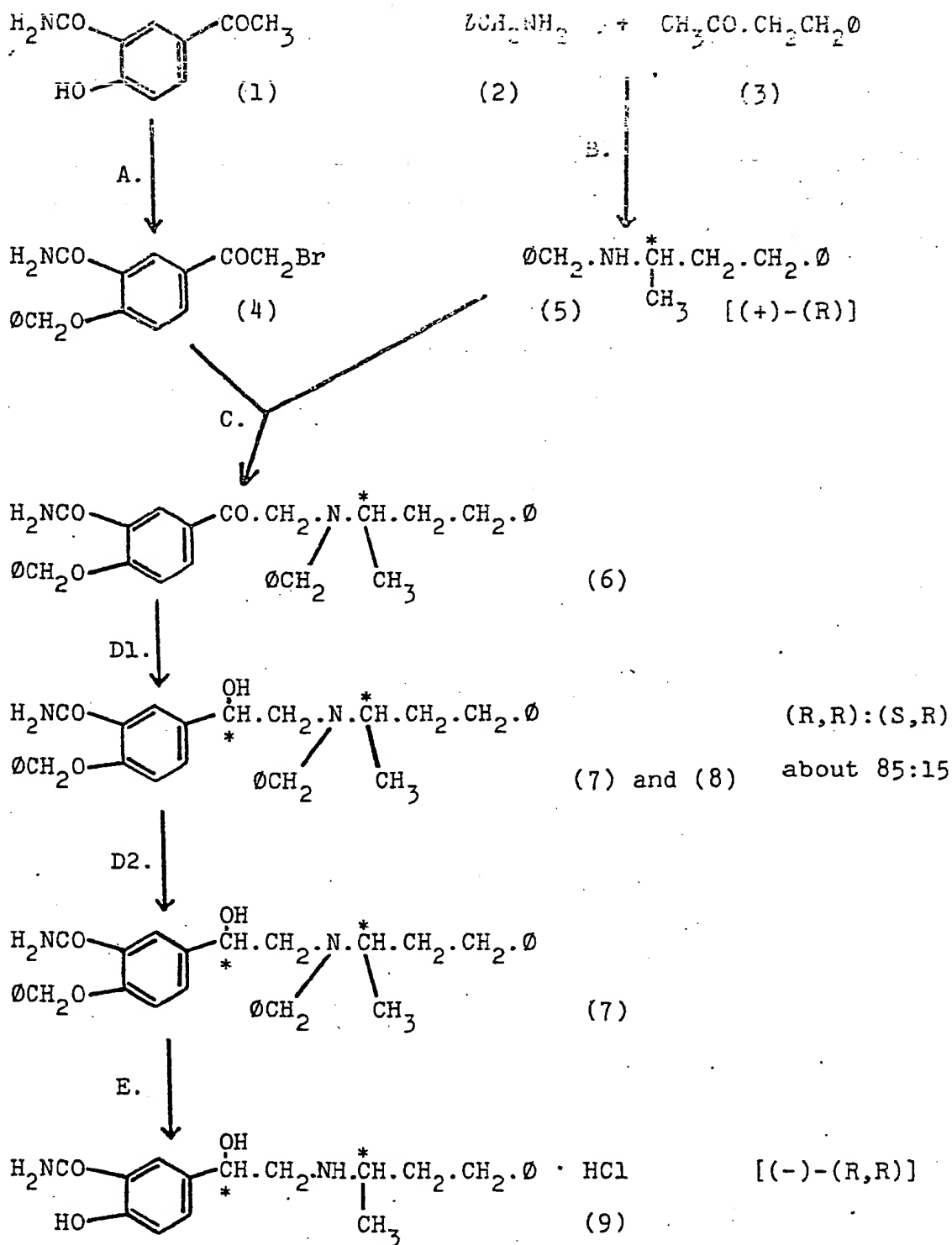
	labetalol			
	1.0; 1.0			
	diastereoisomer A		diastereoisomer B	
	1.2; 0.16		0.5; 1.2	
5	(S,R)-	(R,S)-	(S,S)-	(R,R)-
	isomer	isomer	isomer	isomer
	4.9; 0.05	0.4; 0.3	0.8; 0.02	0.3; 12.3

This table clearly shows the unexpectedly high β -blocking activity and ratio of β :- α -blocking activities possessed by the (R,R)-optical isomer. Additionally, the (R,R)-optical isomer has been found to possess greater direct peripheral vasodilation activity than labetalol, and this also contributes to its anti-hypertensive activity. Moreover, the (R,R)-optical isomer is substantially non-toxic at therapeutic doses.

According to the invention therefore we provide the (R,R)-optical isomer of labetalol, namely 5- {(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl} salicylamide, which can be characterised by means of its hydrochloride salt which is dimorphic with m.pt.s. of about 133-134°C. and about 192-193.5°C. and an $[\alpha]_D^{26}$ of about -30.6° (conc. 1 mg./ml., ethanol), said (R,R)

optical isomer being substantially free of the corresponding (R,S), (S,R) and (S,S) optical isomers, and the pharmaceutically acceptable salts thereof. Such salts include especially the above-mentioned hydrochloride salt, the sulfate, maleate, tartrate and citrate salts, and also the acetate, phthalate, succinate, lactate, malate, cinnamate, hydrobromide and phosphate salts.

As far as we know, no one has succeeded in resolving labetalol by standard methods and no individual optical isomer of labetalol has yet been disclosed. Indeed, we ourselves have tried several standard methods of resolving labetalol by salt formation with an optically active acid, but all these standard methods have failed. In particular, all our attempts to resolve diastereoisomer B of labetalol using the following reagents have failed: N-acetyl-L-leucine, N-tosyl-L-(+)-glutamic acid, N-tosyl-L-leucine and N-tosyl-D-leucine. We have therefore concluded that attempts to resolve diastereoisomer B by such methods are likely to fail and that stereospecific synthesis most probably offers a more fruitful route to the individual isomers of labetalol and in particular to the (R,R) optical isomer thereof. An especially preferred embodiment of such a stereospecific synthesis is shown in the following reaction scheme:



In this reaction scheme:

compounds (6), (7), (8) and (9) are novel and are features of the invention;

"Ø" shows an unsubstituted phenyl group and asterisks indicate asymmetrically substituted carbon atoms; and

5 the various steps can be described in general terms as follows:

A: (i) Formation of phenolate salt with a strong base in an organic solvent;

10 (ii) introduction of a protecting group on to the phenolic oxygen atom;

(iii) bromination with bromine in an inert organic solvent.

15 B: (i) Condensation to a Schiff's base, e.g. by heating in a water-immiscible organic solvent in the presence of an acid catalyst with azeotropic removal of water;

(ii) reduction of the Schiff's base to a secondary amine;

20 (iii) resolution of this secondary amine into its optical antipodes and isolation of the (+)-(R)-optical isomer.

C: Condensation to a tertiary amine by reaction in an inert organic solvent in the presence of an

acid binding agent.

D1: Reduction.

D2: Separation of isomers, e.g. by chromatography.

5 E: Hydrogenolysis of protecting groups with hydrogen
and a catalyst and isolation of the (R,R) optical
isomer e.g. as an acid addition salt.

The novel and inventive process is of course not restricted
exactly to the sequence of steps given above.

10 According to the invention, one process for the preparation
of the novel (R,R) optical isomer of labetalol and of
its pharmaceutically acceptable acid addition salts, sub-
stantially free of the corresponding (R,S), (S,R) and
(S,S) optical isomers, comprises removing the protecting
groups from an N,O-protected (R,R) optical isomer or acid
15 addition salt thereof, the term "N,O-protected" indicating
that the basic nitrogen atom and the phenolic hydroxy group
are protected, and isolating the resulting (R,R) optical
isomer as the free base or as a pharmaceutically acceptable
acid addition salt.

20 The protecting groups are preferably such groups as can be

removed by hydrogenolysis, e.g. by means of hydrogen and palladium on carbon; examples of such groups include the N- or O-benzyl, N- or O-benzyloxycarbonyl, N- or O-benzhydryl, N-trichloroethoxycarbonyl or N-trityl group. Preferably
5 both protecting groups are benzyl groups. Alternatively, the protecting groups can be such groups as are removable by mild hydrolysis, e.g., an N- or O-benzyloxycarbonyl or N-trifluoroacetyl group.

The N,O-protected (R,R) optical isomer used as starting
10 material in the above process according to the invention is preferably obtained by resolution of an N,O-protected (R,R) (S,R) diastereoisomeric mixture. This resolution can conveniently be effected by physical methods, e.g. by chromatography especially on silica gel. This N,O-protected (R,R) (S,R) diastereoisomeric mixture can be obtained
15 by reduction of an N,O-protected 5- { N-[(R)-1-methyl-3-phenylpropyl]glycyl } salicylamide, preferably by means of a borohydride in an organic solvent, especially sodium borohydride in a lower alkanol such as methanol or ethanol.
20 Other reducing agents that can conveniently be used include lithium borohydride or an alkali metal alkylborohydride, e.g. lithium or potassium tri-s-butylborohydride or lithium or potassium tri-(3-methyl-2-butyl)borohydride, in an organic solvent such as tetrahydrofuran.

The N,O-protected 5-[2-(1-methyl-3-phenylpropyl)glycyl] salicylamide can be obtained by condensation of a 4-O-protected- α -bromo-3-carbamoylacetophenone with an (R)-N-protected-1-methyl-3-phenylpropylamine, for example
5 in the presence of an excess of the latter reagent, or of triethylamine or 2,2,6,6-tetramethylpiperidine or especially of potassium carbonate as acid binding agent, and also of dimethylformamide as organic solvent.

10 The 4-O-protected- α -bromo-3-carbamoylacetophenone can be prepared by protection of the phenolic hydroxy group of 5-acetylsalicylamide, e.g. by formation of an alkali salt and benzylation with benzylchloride; the bromine atom in the side chain can then be introduced by bromination with bromine in chloroform.

15 The (R)-N-protected-1-methyl-3-phenylpropylamine can be prepared by condensation of an amine carrying a suitable protecting group, e.g. benzylamine, with benzyl acetone, preferably under reflux in a water-immiscible organic solvent and in the presence of a strong acid catalyst e.g.
20 p-toluenesulfonic acid, with continuous removal of water. The resulting Schiff's base is then reduced under mild conditions, e.g. with sodium borohydride in methanol, so as not to remove the protecting group. Alternatively, the N-protecting group can be directly introduced
25 into 1-methyl-3-phenylpropylamine. The N-protected

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- 1-methyl-3-phenylpropylamine can then be resolved and the (+)-(R)-optical isomer isolated, e.g. with N-p-toluenesulfonyl-(L)-leucine followed by N-acetyl-(L)-leucine, or with N-p-toluenesulfonyl-(D)-leucine. Other reagents that may be used include 2,3,5,6-di-O-isopropylidene-2-keto-(L)-gulonic acid and the D- and L- forms of dibenzoyl tartaric acid, ditoluoyl tartaric acid and mandelic acid. If desired, the resolution can be deferred until step C (condensation) has been carried out.
- 10 A further feature of the present invention comprises a process for the preparation of the (R,R) optical isomer of labetalol and its pharmaceutically acceptable acid addition salts, substantially free of the corresponding (R,S), (S,R) and (S,S) optical isomers, consisting of
- 15 the steps of condensing a 4-O-protected- α -bromo-3-carbamoylacetophenone with an N-protected-1-methyl-3-phenylpropylamine, obtaining an N,O-protected 5- $\{$ N-[(R)-1-methyl-3-phenylpropyl]glycyl $\}$ salicylamide either by using an (R)-N-protected-1-methyl-3-phenylpropylamine
- 20 in the foregoing condensation or by resolving the resulting racemic N,O-protected 5-[N-1-methyl-3-phenylpropyl]glycyl] salicylamide, reducing the N,O-protected 5- $\{$ N-[(R)-1-methyl-3-phenylpropyl]glycyl $\}$ salicylamide to a mixture of an N,O-protected 5- $\{$ (R)-1-hydroxy-2-[(R)-(1-methyl-

-3-phenylpropyl)amino]ethyl} salicylamide and the corresponding (S,R) optical isomer, separating from this mixture the N,O-protected 5- { (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl } salicylamide, removing the
5 protecting groups therefrom and isolating 5- { (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl } salicylamide as the free base or as a pharmaceutically acceptable acid addition salt thereof.

A preferred feature of this process comprises the
10 steps of condensing 4-benzyloxy- α -bromo-3-carbamoyl-acetophenone with (R)-(+)-N-benzyl-1-methyl-3-phenylpropylamine to yield 2-O-benzyl-5- { N-benzyl-N-[(R)-1-methyl-3-phenylpropyl]glycyl } salicylamide, reducing this compound to a mixture of 2-O-benzyl-5- { (R)-1-hydroxy-
15 -2-[(R)-(1-methyl-3-phenylpropyl)benzylamino]ethyl } salicylamide and the corresponding (S,R) optical isomer, separating from this mixture 2-O-benzyl-5- { (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)benzylamino]ethyl } salicylamide, removing the protecting N- and O-benzyl groups
20 therefrom by hydrogenolysis and isolating 5- { (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl } salicylamide as the free base or as a pharmaceutically acceptable acid addition salt thereof.

This preferred process can be carried out under conditions described above. Thus the step of condensing is preferably effected in the presence of potassium carbonate as acid binding agent and of dimethylformamide as inert organic solvent; the step of reducing is preferably effected by means of an alkali metal borohydride in an inert organic solvent; and the step of separating is conveniently effected by chromatography.

The R,R-isomer can be isolated as the free base or as a pharmaceutically acceptable acid addition salt. These salts can be prepared by standard methods.

The hydrochloride salt of the novel (R,R)-optical isomer is dimorphic and exists in two crystalline forms m.pt. about 133-134°C. and about 192-193.5°C. The characterising constants given in the Example (part E) are for the higher melting (presumably the thermodynamically more stable) crystalline form, although either form may be obtained.

The α - and β -blocking activities discussed earlier in this specification can be determined by known methods, e.g. by those of Farmer et.al., Brit. J. Pharm., 45, 660 (1972); Robson, J. Pharm. Exp. Therap., 175, 157 (1970), and Levy, Arch. Int. Pharmacodyn. Ther., 204, 143 (1973).

The following Example describes the preparation of the active compound of the present invention and intermediates in its preparation.

EXAMPLE

5 A. 4-Benzyloxy- α -bromo-3-carbamoylacetophenone (4)

To a solution of 115.4 g. (0.644 mol.) of 5-acetylsalicylamide (1) in 1.2 liter of dimethylformamide add 33.1 g. (0.613 mol.) of sodium methoxide in small portions with cooling and stirring. Heat the mixture on a steam bath and add 75 ml. (0.652 mol.) of benzylchloride dropwise. Continue heating and stirring for 7 hours. After stirring and cooling, pour the mixture into 6 liters of ice-water containing 15 g. of sodium carbonate. Filter, wash well with water, digest with 700 ml. of ethanol, chill and refilter to obtain analytically pure 4-benzyloxy-3-carbamoylacetophenone, m.p. 157-160°C.

To a refluxing, stirred solution of 127.0 g. (0.47 mol.) of 4-benzyloxy-3-carbamoylacetophenone in 1.2 liter of chloroform, add a few ml. of a solution of 76.5 g. (0.49 mol.) bromine in 220 ml. of chloroform and wait until the colour is discharged (ca. 5-10 min.). Cool the hot solution to room temperature and add the remaining bromine

solution dropwise with stirring at room temperature until precipitation begins; then reflux the reaction mixture and continue the dropwise addition. After refluxing for 10 min. following completion of the addition, chill the solution in an ice bath, filter off the solid and wash it with cold chloroform. Stir the crude solid for 20 min. in 800 ml. of ice-cold water, filter it off, wash it well with water and dry it. Recrystallize it from methylethyl ketone to afford two crops of the product (4), m.p. 150-152°C. and m.p. 146-149°C., both of which are usable for the preparation of 2-O-benzyl-5-{ N-benzyl-N-[(R)-1-methyl-3-phenylpropyl]glycyl } salicylamide (6).

B. (R)-(+)-N-Benzyl-1-methyl-3-phenylpropylamine (5)

In an apparatus fitted with a Dean and Stark trap, reflux a solution of 1.0 kg. (6.75 mol.) of benzylacetone (3), 725 g. (6.75 mol.) of benzylamine (2) and 5.0 g. of p-toluenesulfonic acid hydrate in 7 liters of benzene for 14 hours. Remove the solvent in vacuo and dissolve the residue in 6.5 liters of methanol. With cooling and stirring, carefully add 125 g. of sodium borohydride and stir the mixture for 16 hours at room temperature. Remove the methanol in vacuo, add 2 liters of water and 4 liters of benzene, and extract the product

into the benzene. Dry the solution over anhydrous magnesium sulfate, filter, and distil the filtrate, collecting the fraction b.p. 145-150°C/0.5mm. Dissolve 1,028 g. (4.288 mol.) of the distillate and 1,230 g. (4.328 mol.) of N-p-toluenesulfonyl-(L)-leucine in 7.2 liters of boiling ethanol and allow to cool to room temperature without agitation. Wash the resulting precipitate with a small amount of ice-cold ethanol, recrystallize it from 4.8 liters of ethanol, filter off the solid product and wash it with ice-cold ethanol. This solid product is highly enriched with the salt of the undesired (S)-enantiomer. Combine the mother liquors from the original precipitation and from the recrystallization, remove the solvent and recover the free base by basifying with 500 ml of 20% aqueous sodium hydroxide and extracting with benzene. After drying over anhydrous magnesium sulfate, filtering and removing the benzene, dissolve the residue [487 g. (2.04 mol.)] and 346 g. (2.06 mol.) of N-acetyl-(L)-leucine in 2.0 liters of boiling ethanol and allow the solution to cool to room temperature. Filter off the product and recrystallize it once from 1.8 liter of ethanol and then from 4.0 liters of acetonitrile to obtain the salt of the desired (R)-enantiomer with N-acetyl-(L)-leucine, m.p. 151-152°C. Basify with 400 ml. of aqueous 2.5N sodium hydroxide, extract with ether, dry over anhydrous magnesium sulfate, filter

and remove the solvent in vacuo to obtain the product

(5), $[\alpha]_D^{26} = +4.5^\circ$ (c = 5.0, ethanol):

C. 2-O-Benzyl-5-{ N-benzyl-N-[(R)-1-methyl-3-phenyl-propyl]glycyl } salicylamide (6)

5 Stir a mixture of 224 g. (0.94 mol.) of (R)-(+)-N-benzyl-1-methyl-3-phenylpropylamine (5), 372 g. (ca. 1.07 mol.) of 4-benzyloxy- α -bromo-3-carbamoylacetophenone (4) and 372 g. (2.7 mol.) of potassium carbonate in 1.6 liters dimethylformamide at room temperature for 4 hours (reaction mildly
10 exothermic). Add 8.7 liters water and extract with ether, dry over anhydrous sodium sulfate, filter, and remove the ether in vacuo (up to 30-40°C.) to yield the crude product (6) as a syrup.

D. 2-O-Benzyl-5-{ (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl) benzylamino]ethyl } salicylamide (7)
15

1.(a) Dissolve 520 g. (not more than 0.94 mol.) of crude 2-O-benzyl-5-{ N-benzyl-N-[(R)-1-methyl-3-phenylpropyl]glycyl } salicylamide (6) in 3.1 liters ethanol and, with stirring and cooling, add portionwise 35.5 g. (0.94 mol.)
20 of sodium borohydride. Stir the mixture at room temperature for 16 hours, remove the solvent in vacuo, add 3.2 liters water and heat the mixture for 30 minutes on a steam

bath. Cool, extract with benzene, dry the benzene layer over anhydrous magnesium sulfate, filter, remove the solvent in vacuo, and obtain the crude product as a syrup (ratio R,R:S,R ca. 85:15).

5 The following procedures show that this reduction can be carried out by a variety of further reducing agents:

10 (b) Add one ml. of 0.5M potassium tri-sec-butylborohydride solution in tetrahydrofuran to a cold solution of 200 mg. of the crude aminoketone (6) from step C in 10 ml. of tetrahydrofuran with stirring and cooling in an ice bath. Stir for another thirty minutes. Heat a sample with a few drops of methanol; thin layer chromatography on silica gel using chloroform:ethyl acetate (3:1) as developing solvent shows that the ratio of isomers (R,R):(S,R) in the product is about 70:30.

15 (c) Add 0.5 ml. of 1.0M lithium tri-sec-butylborohydride solution in tetrahydrofuran to a cold solution of 100 mg. of the crude aminoketone (6) from step C in 10 ml. of tetrahydrofuran with stirring and cooling in an ice bath. Stir for another ten minutes, then hydrolyse with a few drops of ethanol. Distil off the solvent and extract the residue with benzene and water. Separate the benzene layer, dry it over anhydrous sodium sulfate, filter and distil off the benzene. The ratio of isomers (R,R):(S,R) in the product is shown by pmr spectrum to be about 80:20.

20

25

(d) Replacement of the solvent (10 ml. of tetrahydrofuran) for the crude aminoketone (6) with 10 ml. of benzene in procedures (b) and (c) gives almost identical results to those of procedures (b) and (c).

5 (e) Dissolve 63.0 g. of crude aminoketone (6) from step C in a mixture of 960 ml. of benzene and 40 ml. of tetrahydrofuran, and decant off the solution from insoluble material. Add this solution dropwise to a suspension of 4.20 g. of lithium borohydride in 480 ml. of benzene and 20 ml. of tetrahydrofuran with stirring and cooling in an ice-bath. Stir for an additional two hours, then decompose with water. Separate the organic layer and dry it over anhydrous sodium sulfate. Filter and evaporate the filtrate to dryness on a rotary evaporator. The ratio of isomers (R,R):(S,R) in the product is shown by pmr spectrum to be about 70:30.

20 2.(a) Chromatograph 47 g. of the crude mixture from e.g. 1.(a) above on 1.5 kg. of thin layer grade silica gel with chloroform : ethyl acetate (3:1) and obtain the pure product (7), which is eluted first. This compound's pmr spectrum in CDCl_3 is assigned as follows: $\delta = 1.02$ (d, C-CH_3 ; $J = 7$ Hz), 1.42-2.00 (m, $-\text{C}(\text{CH}_3)-\text{CH}_2-$), 3.66 (q, $\text{N-CH}_2-\text{C}_6\text{H}_5$), 4.62 (q, CHOH), 5.15 (s, $\text{OCH}_2-\text{C}_6\text{H}_5$).

(b) The product from 1(b) to (e) can similarly be resolved: e.g., chromatography of the product (53 g.) from 1(e) on 1,200 g. of a silica gel column using chloroform:ethyl-acetate (3:1) provides the pure product (7).

5 E. (-)-5-[(R)-1-Hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)-amino]ethyl] salicylamide hydrochloride salt (9)

Treat a solution of 3.0 g. (0.0059 mol.) of 2-O-benzyl-5-[(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)benzylamino]ethyl] salicylamide in 30 ml. of ethyl ether with 2N ethereal hydrogen chloride until no further precipitation occurs. Wash the precipitated 2-O-benzyl-5-[(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)benzylamino]ethyl] salicylamide hydrochloride with ether to remove excess hydrogen chloride and dissolve it in 100 ml. ethanol. To the ethanol solution add 300 mg. of a 20% palladium hydroxide on carbon catalyst and hydrogenate (3 atm.; 3.1 kg. cm.⁻²) in a Paar apparatus with shaking at room temperature for 3 hours. Filter off the catalyst, evaporate, and triturate the solid residue with isopropanol. Dissolve the solid in 11 ml. of 1N sodium hydroxide, adjust the pH to about 8 and precipitate the free base by bubbling in carbon dioxide. Collect the free base, wash it with water and dry it in vacuo at 40°C. Chromatograph the free base on 450 g. of silica gel and dissolve the pure product in 20 ml. of boiling acetonitrile. Cool the solution and carefully acidify with 2N ethereal HCl to about pH2. Solidify the gum which precipitates by refluxing the mixture for 10 minutes, filter off the solid, wash it with ethyl ether and recrystallize it from ethanol to obtain analytically pure product (9), m.p. 192-193.5°C.(dec.), $[\alpha]_D^{26} = -30.6^\circ$ (c=1.0, ethanol).

30 The (R,R) isomer and its pharmaceutically acceptable salts are useful in the treatment of cardiovascular disorders and particularly in the treatment of mammalian hypertension. They can also be used for direct peripheral vasodilation and in the treatment of glaucoma. They are preferably

administered orally but can also be administered by injection. Laboratory tests indicate that the effective oral dose (ED_{50}) for the (R,R) isomer or a pharmaceutically acceptable salt thereof will typically lie within the range of 0.01 to 5 25 mg./kg., preferably 0.5 to 5 mg./kg., of mammalian weight.

The invention therefore provides pharmaceutical compositions containing as active ingredient 5- $\left\{ \begin{array}{l} \text{(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl} \end{array} \right\}$ salicylamide or a pharmaceutically acceptable acid addition salt thereof, together with a pharmaceutical carrier or excipient, said (R,R) optical isomer being substantially free of the corresponding (R,S), (S,R) and (S,S) optical isomers. The compositions are preferably in the form of 10 dosage units, e.g. tablets, pills, capsules, suppositories or injectable preparations in ampoules. The compositions may also be for example in the form of powders, syrups, elixirs or suspensions. The required daily dosage may be administered in single or divided doses. The exact dose to be 20 administered will, of course, be dependent upon various factors such as the age and weight of the subject mammal and the individual response. Dosage units preferably contain from 2 to 500 mg., more preferably from 10 to 200 mg., of the (R,R) isomer according to the invention (or pharmaceutically acceptable acid addition salt thereof). 25

Typical pharmaceutically acceptable carriers for use in formulations described above are exemplified by: sugars such as lactose, sucrose, mannitol and sorbitol; starches such as corn starch, tapioca starch and potato starch; 5 cellulose and derivatives such as sodium carboxymethyl cellulose, ethyl cellulose and methyl cellulose; calcium phosphates such as dicalcium phosphate and tricalcium phosphate; sodium sulfate; calcium sulfate; polyvinylpyrrolidone; polyvinyl alcohol; stearic acid; 10 alkaline earth metal stearates such as magnesium stearate and calcium stearate; vegetable oils such as peanut oil, cottonseed oil, sesame oil, olive oil and corn oil; non-ionic, cationic and anionic surfactants; ethylene glycol polymers; β -cyclodextrin; fatty 15 alcohols; hydrolyzed cereal solids; and other non-toxic compatible fillers, binders, disintegrants, and lubricants commonly used in pharmaceutical formulations.

In treating certain patients with the R,R-isomer of this invention, it may be desirable to include other pharmaceutically active ingredients in the same composition. 20 For example, in treating patients in whom salt and water retention is a problem, effective amounts of a diuretic, e.g., hydrochlorothiazide or trichloromethiazide, may be included.

Pharmaceutical Formulations

In the following examples, the active ingredient is preferably 5- $\left\{ \text{(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenyl-propyl)amino]ethyl} \right\}$ salicylamide hydrochloride, but an equivalent quantity of the (R,R) isomer itself or of another pharmaceutically acceptable acid addition salt, especially a salt named herein, may be substituted:

<u>Injectable Solution:</u>		<u>mg./ml.</u>
	Active ingredient	5.00
10	Methyl <u>p</u> -hydroxybenzoate	0.80
	Propyl <u>p</u> -hydroxybenzoate	0.10
	Disodium Edetate	0.10
	Citric Acid Monohydrate	0.08
	Dextrose	40.00
15	Water for injection qs ad	1.00 ml.

Manufacturing Procedure:

Dissolve the p-hydroxybenzoates in a portion of water for injection at 60-70°C., and cool the solution to 25-35°C. Charge and dissolve all other excipients and the active ingredient. Bring the solution to final volume, filter it through a sterilizing membrane and fill into sterile containers.

Oral Formulations:a) Capsules:

	Formula	Quantities per capsule	
		(mg.)	(mg.)
	Active ingredient	200.0	100.0
5	Lactose	223.0	111.5
	Corn Starch	75.0	37.5
	Magnesium Stearate	<u>2.0</u>	<u>1.0</u>
		500.0	250.0

Manufacturing Procedure:

- Blend the active ingredient, lactose and corn starch
 10 until uniform; then blend the magnesium stearate
 into the resulting powder. Encapsulate the mixture
 into suitably sized two-piece hard gelatin capsules.

b) Tablets

	Formula	Quantities per tablet	
		(mg.)	(mg.)
15	Active ingredient	200.0	100.0
	Lactose	211.0	105.5
	Corn Starch	12.0	6.0
	Water (per thousand tablets)	(120 ml.)*	(60 ml.)*
	Corn Starch	75.0	37.5
20	Magnesium Stearate	<u>2.0</u>	<u>1.0</u>
		500.0	250.0

* (The water evaporates during manufacture.)

Manufacturing Procedure:

Blend the active ingredient with the lactose until uniform. Blend the smaller quantity of corn starch with the water and add the resulting corn starch paste, then mix
5 until a uniform wet mass is formed. Add the remaining corn starch to the resulting wet mass and mix until uniform granules are obtained. Screen the granules through a suitable milling machine, using a 3/4" stainless steel screen. Dry the milled granules in a suitable drying oven
10 until the desired moisture content is obtained. Mill the dried granules through a suitable milling machine using 16 mesh stainless steel screen. Blend in the magnesium stearate and compress the resulting mixture
15 into tablets of desired shape, thickness, hardness and disintegration.

1. The (R,R) optical isomer of labetalol, namely 5-
 { (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl }
 salicylamide, said (R,R) optical isomer being substan-
 tially free of the corresponding (R,S), (S,R) and (S,S) op-
 tical isomers; and the pharmaceutically acceptable acid addi-
 tion salts thereof, especially its hydro-
 choride, sulfate, maleate, tartrate, citrate, acetate, phtha-
 late, succinate, lactate, malate, cinnamate, hydrobromide
 and phosphate.

2. 5-[(R)-1-Hydroxy-2-[(R)-(1-methyl-3-phenylpropyl) amino]ethyl] salicylamide and its hydrochloride, said hydrochloride being a dimorphic compound with m.pt.s. of about 133-134°C. and about 192-193.5°C., and $[\alpha]_D^{26}$ of about -30.6° (conc. 1 mg./ml., ethanol); said (R,R) optical isomer being substantially free of the corresponding (R,S), (S,R) and (S,S) optical isomers.

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the phenolic hydroxy group are protected, and isolating the resulting (R,R) optical isomer as the free base or as a pharmaceutically acceptable acid addition salt.

4. A process as claimed in claim 3 wherein the step of removing the protecting groups is performed by hydrogenolysis, preferably by means of hydrogen and palladium on carbon, of protecting groups selected from N- or O-benzyl, N- or O-benzyloxycarbonyl or N-trichloroethoxycarbonyl groups, in particular of N- and O-benzyl groups.

10 5. A process as claimed in claim 3 or claim 4 wherein the N,O-protected (R,R) optical isomer has been obtained by resolution of an N,O-protected (R,R)-(S,R) diastereoisomeric mixture, by physical methods, in particular by chromatography on silica gel.

15 6. A process as claimed in claim 5 wherein the N,O-protected (R,R)-(S,R) diastereoisomeric mixture has been obtained by reduction of an N,O-protected 5- $\left\{ \text{N}-[(\text{R})\text{-1-methyl-3-phenylpropyl}]\text{glycyl} \right\}$ salicylamide, preferably by a borohydride in an organic solvent, in particular by sodium
20 borohydride in a lower alkanol.

7. A process as claimed in claim 6 wherein the N,O-protected 5- $\left\{ \text{N}-[(\text{R})\text{-1-methyl-3-phenylpropyl}]\text{glycyl} \right\}$ salicylamide has been obtained by condensation of a 4-O-protected α -bromo-3-carbamoylacetophenone with an (R)-N-protected-1-methyl-
25 -3-phenylpropylamine.

8. A process for the preparation of the (R,R) optical isomer claimed in claim 1 and of its pharmaceutically acceptable acid addition salts, substantially free of the corresponding (R,S), (S,R) and (S,S) optical isomers, which comprises condensing a 4-O-protected- α -bromo-3-
5 -carbamoylacetophenone with an N-protected-1-methyl-3-phenylpropylamine, obtaining an N,O-protected 5- $\left\{ \text{N}-[(\text{R})-1\text{-methyl-3-phenylpropyl}] \text{glycyl} \right\}$ salicylamide either by using an (R)-N-protected-1-methyl-3-phenylpropylamine
10 in the foregoing condensation or by resolving the resulting racemic N,O-protected 5-[N-1-methyl-3-phenylpropyl]glycyl] salicylamide, reducing the N,O-protected 5- $\left\{ \text{N}-[(\text{R})-1\text{-methyl-3-phenylpropyl}] \text{glycyl} \right\}$ salicylamide to a mixture of an N,O-protected 5- $\left\{ (\text{R})-1\text{-hydroxy-2}-[(\text{R})-(1\text{-methyl-3-phenylpropyl})\text{amino}] \text{ethyl} \right\}$ salicylamide and the cor-
15 responding (S,R) optical isomer, separating from this mixture the N,O-protected 5- $\left\{ (\text{R})-1\text{-hydroxy-2}-[(\text{R})-(1\text{-methyl-3-phenylpropyl})\text{amino}] \text{ethyl} \right\}$ salicylamide, removing the protecting groups therefrom and isolating 5- $\left\{ (\text{R})-1\text{-hydroxy-2}-[(\text{R})-(1\text{-methyl-3-phenylpropyl})\text{amino}] \text{ethyl} \right\}$ salicylamide as the free base or as a pharmaceutically acceptable acid addition salt thereof; in particular the process comprising condensing 4-benzyloxy- α -bromo-3-carbamoylacetophenone with (R)-(+)-N-benzyl-1-methyl-3-phenyl-

propylamine to yield 2-O-benzyl-5-{N-benzyl-N-[(R)-1-methyl-3-phenylpropyl]glycyl} salicylamide, reducing this compound to a mixture of 2-O-benzyl-5-{(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)benzylamino]ethyl} salicylamide and the corresponding (S,R) optical isomer, separating from this mixture 2-O-benzyl-5-{(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)benzylamino]ethyl} salicylamide, removing the protecting N- and O-benzyl groups therefrom by hydrogenolysis and isolating 5-{(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl} salicylamide as the free base or as a pharmaceutically acceptable acid addition salt thereof.

9. Pharmaceutical compositions containing as active ingredient 5-{(R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl} salicylamide, characterized as having a hydrochloride salt which has a melting point of about 192-193.5°C. and an $[\alpha]_D^{26}$ of about -30.6° (conc. 1 mg./ml., ethanol), said (R,R) optical isomer being substantially free of the corresponding (R,S), (S,R) and (S,S) optical isomers, or a pharmaceutically acceptable acid addition salt thereof, together with a pharmaceutical carrier or excipient.

10. Compositions as claimed in claim 9 in the form of dosage units, e.g., tablets, capsules, pills, suppositories or injectable preparations in ampoules, containing preferably from 2 to 500 mg. of active ingredient per dosage unit.
11. N,O-Protected 5- { (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)amino]ethyl } salicylamides, diastereoisomeric mixtures of this (R,R) optical isomer with its (S,R) diastereoisomer, and N,O-protected 5- { N-[(R)-1-methyl-3-phenylpropyl]glycyl } salicylamides, wherein the basic nitrogen atom and the phenolic hydroxy group are protected by hydrogenolysable or readily hydrolysable groups; in particular the compounds
- 2-O-benzyl-5- { (R)-1-hydroxy-2-[(R)-(1-methyl-3-phenylpropyl)benzylamino]ethyl } salicylamide and 2-O-benzyl-5- { N-benzyl-N-[(R)-1-methyl-3-phenylpropyl]glycyl } salicylamide.



European Patent
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EUROPEAN SEARCH REPORT

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Application number

EP 79 10 3470

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int. Cl. 3)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
D, X	GB - A - 1 266 058 (ALLEN & HAN-BURYS) * Whole patent, particularly page 2, lines 38-46 *	1-11	C 07 C 103/29 A 61 K 31/165// C 07 C 87/28
D, X	DE - A - 2 616 403 (SCHERICO) * Whole patent *	1, 2, 9, 10	TECHNICAL FIELDS SEARCHED (Int. Cl. 3) C 07 C 103/29 A 61 K 31/165
			CATEGORY OF CITED DOCUMENTS X: particularly relevant A: technological background O: non-written disclosure P: intermediate document T: theory or principle underlying the invention E: conflicting application D: document cited in the application L: citation for other reasons
			&: member of the same patent family. corresponding document
The present search report has been drawn up for all claims			
Place of search The Hague		Date of completion of the search 14-1-1979	Examiner ALLARD

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